```
; Sequence 24, Application US/08879338A
; Patent No. 6063906
; GENERAL INFORMATION:
  APPLICANT: Brenner, Michael B.
  APPLICANT: Parker, Christina M.
  TITLE OF INVENTION: Antibodies to No. 6063906el Integrin Alpha
; TITLE OF INVENTION: Subunit
; FILE REFERENCE: B0801/7080/ERP
; CURRENT APPLICATION NUMBER: US/08/879,338A
; CURRENT FILING DATE: 1997-06-20
; EARLIER APPLICATION NUMBER: US 08/663,731
; EARLIER FILING DATE: 1996-06-14
; EARLIER APPLICATION NUMBER: US 08/199,776
; EARLIER FILING DATE: 1994-02-18
; NUMBER OF SEQ ID NOS: 31
  SOFTWARE: FastSEQ for Windows Version 3.0
; SEQ ID NO 24
   LENGTH: 12
   TYPE: PRT
   ORGANISM: Artificial Sequence
   OTHER INFORMATION: Synthetic Peptide
US-08-879-338-24
 Query Match
                          2.0%; Score 8; DB 3; Length 12;
 Best Local Similarity 100.0%; Pred. No. 1.3;
            8; Conservative 0; Mismatches
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Qу
         385 EDEEEEEE 392
              111111
Db
           1 EDEEEEEE 8
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; Sequence 24, Application US/08199776
; Patent No. 5594120
  GENERAL INFORMATION:
     APPLICANT: Brenner, Michael B. APPLICANT: Parker, Christina M.
     TITLE OF INVENTION: No. 5594120el integrin alpha subunit
     NUMBER OF SEQUENCES: 25
    CORRESPONDENCE ADDRESS:
       ADDRESSEE: Wolf, Greenfield and Sacks, P.C.
       STREET: 600 Atlantic Avenue
       CITY: Boston
       STATE: MA
       COUNTRY: USA
       ZIP: 02210
    COMPUTER READABLE FORM:
       MEDIUM TYPE: Floppy disk
       COMPUTER: IBM PC compatible
       OPERATING SYSTEM: PC-DOS/MS-DOS
       SOFTWARE: PatentIn Release #1.0, Version #1.25
    CURRENT APPLICATION DATA:
      APPLICATION NUMBER: US/08/199,776
       FILING DATE:
       CLASSIFICATION: 514
    ATTORNEY/AGENT INFORMATION:
      NAME: Plumer, Elizabeth R.
       REGISTRATION NUMBER: 36,637
       REFERENCE/DOCKET NUMBER: B0801/7020
    TELECOMMUNICATION INFORMATION:
       TELEPHONE: 617-720-3500
       TELEFAX: 617-720-2441
  INFORMATION FOR SEQ ID NO: 24:
   SEQUENCE CHARACTERISTICS:
      LENGTH: 12 amino acids
      TYPE: amino acid
      STRANDEDNESS: single
;
      TOPOLOGY: linear
;
    MOLECULE TYPE: peptide
    HYPOTHETICAL: YES
    ANTI-SENSE: NO
    ORIGINAL SOURCE:
       ORGANISM: synthetic peptide
US-08-199-776-24
 Query Match
                          2.0%; Score 8; DB 1; Length 12;
 Best Local Similarity 100.0%; Pred. No. 1.3;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps
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         385 EDEEEEEE 392
Qу
             Db
           1 EDEEEEEE 8
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ID
     O8TAD8
                PRELIMINARY; PRT; 396 AA.
AC
     Q8TAD8;
     01-JUN-2002 (TrEMBLrel. 21, Created)
DT
     01-JUN-2002 (TrEMBLrel. 21, Last sequence update)
DT
     01-OCT-2002 (TrEMBLrel. 22, Last annotation update)
DT
DΕ
     Smad nuclear interacting protein (Smad nuclear-interacting protein
DE
     1).
GN
     SNIP1.
OS
     Homo sapiens (Human).
OC
     Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC
     Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX
    NCBI TaxID=9606;
RN
     [1]
RP
     SEQUENCE FROM N.A.
RC
     TISSUE=Testis;
RA
     Strausberg R.;
RL
     Submitted (APR-2002) to the EMBL/GenBank/DDBJ databases.
RN
     [2]
RΡ
     SEQUENCE FROM N.A.
RX
    MEDLINE=20347038; PubMed=10887155;
RΑ
    Kim R.H., Wang D., Tsang M., Martin J., Huff C., Caestecker M.P.,
     Parks T.W., Meng X., Lechleider R.J., Wang T., Roberts A.B.;
RΑ
RT
     "A novel smad nuclear interacting protein, SNIP1, suppresses p300-
    dependent TGF-beta signal transduction.";
RT
RT.
    Genes Dev. 14:1605-1616(2000).
RN
RP
    SEQUENCE FROM N.A.
RA
    Lin Y., Martin J., Gruendler C., Mach M., Meng X., Li B.-Y.,
    Lechleider R.J., Huff C., Kim R.H., Grasser W., Paralkar V., Wang T.;
RA
RT
     "Smad1 interaction with antizyme and proteasome beta subunit HsN3 in
RT
    signal transduction of BMPs.";
    Submitted (MAR-2002) to the EMBL/GenBank/DDBJ databases.
RL
DR
    EMBL; BC027040; AAH27040.1; -.
    EMBL; AY081909; AAL91140.1; -.
DR
    InterPro; IPR000253; FHA.
DR
    Pfam; PF00498; FHA; 1.
DR
DR
    SMART; SM00240; FHA; 1.
    PROSITE; PS50006; FHA DOMAIN; 1.
DR
SO
    SEQUENCE 396 AA; 45777 MW; B183F83EC3184676 CRC64;
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                        100.0%; Score 396; DB 4; Length 396;
                        100.0%; Pred. No. 0;
 Best Local Similarity
 Matches 396; Conservative
                             0; Mismatches
                                               0; Indels
                                                            0; Gaps
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QУ
             Db
           1 MKAVKSERERGSRRRHRDGDVVLPAGVVVKQERLSPEVAPPAHRRPDHSGGSPSPPTSEP 60
Qу
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             Db
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Qv
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AC
     AAG93323;
XX
DT
     13-SEP-2001 (first entry)
XX
DΕ
     Human protein HP10101.
XX
     Human; gene therapy; tumour.
ΚW
XX
OS
     Homo sapiens.
XX
PN
     W0200142302-A1.
XX
PD
     14-JUN-2001.
XX
PF
     06-DEC-2000; 2000WO-JP08631.
XX
PR
     06-DEC-1999;
                    99JP-0346863.
                                            Late 69/890,688
PR
     06-DEC-1999;
                    99JP-0346864.
PR
     08-FEB-2000; 2000JP-0031062.
     10-FEB-2000; 2000JP-0034090.
                                                         2003014475 Al
     10-FEB-2000; 2000JP-0034091.
PR
     14-FEB-2000; 2000JP-0035829.
PR
     14-FEB-2000; 2000JP-0035899.
PR
PR
     14-MAR-2000; 2000JP-0071161.
PR
     30-MAY-2000; 2000JP-0160851.
XX
PΑ
     (NISC-) JAPAN SCI & TECHNOLOGY CORP.
XX
PΙ
     Kato S, Eguchi C, Saeki M;
XX
DR
     WPI; 2001-381646/40.
DR
     N-PSDB; AAH68608.
XX
PT
     Human protein originated from tumor cell line, applicable as drug,
PT
     reagent for studying intracellular protein networks and protein source
PT
     for drug screening, also encoded cDNA for gene diagnosis and gene
PΤ
     therapy -
XX
XX
CC
     The present sequence is a human protein. The human protein, preferably
CC
     originated from tumour cell line, is applicable as a drug, a reagent for
CC
     studying intracellular protein networks and a protein source for
CC
     screening proteins for binding low molecular weight drugs. The human
CC
     protein coding sequence is useful for gene diagnosis and gene therapy,
CC
     expression vectors and transformant cells for detection of ligands and
CC
     receptors.
XX
SQ
     Sequence
                396 AA;
 Query Match
                          100.0%; Score 396; DB 22; Length 396;
 Best Local Similarity
                         100.0%; Pred. No. 0;
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XX
DT
     30-MAR-2001 (first entry)
XX
DE
     Human transcriptional regulator protein #28.
XX
KW
     Human; transcriptional regulator protein; TXREG.
XX
OS
     Homo sapiens.
XX
PN
     W0200078954-A2.
XX
PD
     28-DEC-2000.
XX
PF
     15-JUN-2000; 2000WO-US16766.
XX
PR
     18-JUN-1999;
                  99US-0140109.
XX
PΑ
     (INCY-) INCYTE GENOMICS INC.
XX
PΙ
     Lal P, Yue H, Tang YT, Baughn MR, Azimzai Y, Tran B;
XX
     WPI; 2001-041425/05.
DR
XX
РΤ
     Isolated polypeptide with a human transcriptional regulator protein
PT
     sequence is useful for the diagnosis, prevention and treatment of
PT
     disorders associated with the immune, reproductive and cardiovascular
PT
     systems -
XX
PS
     Claim 1; Page 117-118; 142pp; English.
XX
CC
    The present invention relates to human transcriptional regulator
CC
    protein (TXREG) sequences. The antagonist and an agonist of the proteins
    of the invention are used to treat disorders associated with decreased
CC
CC
    or increased expression or activity of TXREG.
XX
SO
    Sequence
               396 AA;
 Query Match
                         100.0%; Score 396; DB 22; Length 396;
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AC

AAB61328;